

MEDIS SODIUM BICARBONATE- sodium bicarbonate powder
Humco Holding Group, Inc

Medis Sodium Bicarbonate

Drug Facts

Active Ingredient

Sodium Bicarbonate, USP

Purpose

Antacid

Use

For relief of heartburn, acid indigestion, and upset stomach associated with these symptoms.

Warnings

NOT FOR INJECTIONS

Except under supervision of a doctor do not administer to children under 6 years of age.

Do not take more than six, 1/2 tsp. per person up to 60 years old, or three 1/2 tsp. per person 60 years or older in a 24 hour period.

Do not use this product if you are on a sodium restricted diet (each 1/2 tsp. contains 30 mEq (0.7 g) Sodium).

Do not use the maximum dose more than 2 weeks.

Ask a doctor or pharmacist before use if

you are currently taking a prescription drug. Antacids may interact with certain prescription drugs.

Keep out of reach of children.

In case of accidental overdose, seek professional assistance or contact a Poison Control Center immediately.

Directions

Adult and children 6 yrs of age and older:

Take 1/2 tsp. in 1/2 g;ass (4 fl oz) of water every 2 hrs. up to maximum dosage or as directed by doctor.

Inactive ingredients

none

Label



**BICARBONATO DE SODA
POLVO INGERIBLE U.S.P.**

**Sodium Bicarbonate Oral Powder U.S.P.
(BICARBONATE OF SODA) U.S.P. AND FOOD GRADE**

NOT FOR MAKING INJECTIONS.

USE: Anti-Acid, and for baking.
INDICATIONS: For the relief of Heartburn, Sour Stomach, and/or Acid Indigestion.
DOSE: 0.3 to 2 grams 1 to 4 times daily. Should not be given to young children (to age six), unless prescribed by their Physician.
WARNING: KEEP OUT THE REACH OF CHILDREN. In case of accidental overdose, seek professional assistance or contact a Poison Control Center immediately.

**CONTENTS 4 OUNCES (113g.)
DIST. BY HUMCO, TEXARKANA, TEXAS 75501**

USP: Antídoto y para cocinar.
Indicaciones: Para el alivio de Acidez Estomacal No Aguda. Dosis: 0.3 a 2 gramos 1 a 4 veces diarias. No administrar a niños menores de 6 años, excepto por orden médica.
ADVERTENCIA: Manténgase fuera del alcance de los niños. En caso de sobredosis accidental, consulte a un médico o póngase en contacto inmediatamente con un Centro de Control de Envenenamiento.
ADVERTENCIA: No se administre más de 2 gramos en un período de 24 horas, o use la dosis máxima de este producto por más de 2 semanas excepto bajo indicaciones y vigilancia médica.
ADVERTENCIA: Use este producto solamente bajo vigilancia médica si usted está en una dieta restringida de sodio. Contiene 270 mg. de sodio por gramo. Como con cualquier medicamento, si usted está embarazada o está amamantando un bebé, consulte a un médico o profesional de la salud antes de usar este producto.

NO USAR PARA INYECCIONES
 Bicarbonato De Soda Polvo Ingerible U.S.P.

SELO DE SEGURIDAD ADVERTENCIA. No usar si la Banda de Seguridad impresa "Sealed For Your Protection" está rota o falta. Este producto es sellado con una banda alrededor de la tapa o con una hoja delgada de metal bajo la tapa.

WARNING: Do not use if Tamper Evident Seal imprinted "Sealed For Your Protection" is broken or missing. This product is sealed with either a shrink band around cap or foil seal under cap.
WARNING: Do not take more than 2 grams in a 24 hour period, or use the maximum dosage of this product for more than 2 weeks, except under the advice and supervision of a physician.
WARNING: Do not use this product except under the advice of a Physician if you are on a Sodium Restricted Diet. Contains 270 mg. Sodium per Gram. As with any drug, if you are pregnant or nursing a baby, seek the advice of a health professional before using this product.

0395047670 4

R071805RLS

MEDIS SODIUM BICARBONATE

sodium bicarbonate powder

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:0802-2685
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
SODIUM BICARBONATE (UNII: 8MDF5V39QO) (SODIUM CATION - UNII:LYR4M0NH37)	SODIUM BICARBONATE	1000 mg in 1 g

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0802-2685-94	113 g in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	11/09/2017	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M001	11/09/2017	

Labeler - Humco Holding Group, Inc (825672884)

Registrant - Pharma Nobis, LLC (118564114)

Establishment

Name	Address	ID/FEI	Business Operations
Pharma Nobis, LLC		118564114	label(0802-2685) , manufacture(0802-2685) , pack(0802-2685) , analysis(0802-2685)

Revised: 12/2023

Humco Holding Group, Inc